

**Amendments to the Claims:**

Claims 1-4 (Cancelled).

5. (Currently Amended) A compound ~~according to claim 4~~ selected from the group consisting of:

3-{4-Cyano-3-[(R)-2-hydroxy-3-(2-indan-2-yl-1,1-dimethyl-ethylamino)-propoxy]-phenyl}-propionic ethyl ester ;

3-{4-Cyano-3-[(R)-2-hydroxy-3-(2-indan-2-yl-1,1-dimethyl-ethylamino)-propoxy]-phenyl}-propionic acid ;

3-{4-Cyano-3-[(R)-2-hydroxy-3-(2-indan-2-yl-1,1-dimethyl-ethylamino)-propoxyl]-phenyl}-propionic acid isopropyl ester ;

3-{4-Cyano-3-[(R)-2-hydroxy-3-(2-indan-2-yl-1,1-dimethyl-ethylamino)-propoxy]-phenyl}-propionic acid 2-ethoxy ethyl ester ;

3-{4-cyano-3-[(R)-2-hydroxy-3-(2-indan-2-yl-dimethyl-ethylamino)-propoxy]-phenyl}-propionic acid 2-methoxy-1-methyl-ethyl ester ;

3-(4-Cyano-3-[(R)-3-[1,1-dimethyl-2-(5,6,7,8-tetrahydro-naphthalen-2-yl)-ethylamino]-2-hydroxy-propoxy]-phenyl)-propionic acid ;

3-(4-Cyano-3-[(R)-3-[1,1-dimethyl-2-(5,6,7,8-tetrahydro-naphthalen-2-yl)-ethylamino]-2-hydroxy-propoxy]-phenyl)-propionic acid ethyl ester ;

3-(3-Cyano-4-[(R)-3-[1,1-dimethyl-2-(5,6,7,8-tetrahydro-naphthalen-2-yl)-ethylamino]-2-hydroxy-propoxy]-phenyl)-propionic acid ;

3-(3-Cyano-4-[(R)-3-[1,1-dimethyl-2-(5,6,7,8-tetrahydro-naphthalen-2-yl)-ethylamino]-2-hydroxy-propoxy]-phenyl)-propionic acid ethyl ester ;

3-{4-Cyano-3-[(R)-2-hydroxy-3-(2-indan-5-yl-1,1-dimethyl-ethylamino)-propoxy]-phenyl}-propionic acid ; and

3-{4-Cyano-3-[(R)-2-hydroxy-3-(2-indan-5-yl-1,1-dimethyl-ethylamino)-propoxy]-phenyl}-propionate ethyl ester; and pharmaceutically acceptable salts and complexes thereof.

6. (Original) A compound according to claim 5 selected from the group consisting of:

3-{4-Cyano-3-[(R)-2-hydroxy-3-(2-indan-2-yl-1,1-dimethyl-ethylamino)-propoxy]-phenyl}-propionic ethyl ester ; and

3-{4-Cyano-3-[(R)-2-hydroxy-3-(2-indan-2-yl-1,1-dimethyl-ethylamino)-propoxy]-phenyl}-propionic acid ;

and pharmaceutically acceptable salts and complexes thereof.

Claims 7-8 (Cancelled).

9. (Currently Amended) A method ~~according to claim 8 wherein the of treating a bone or mineral disease or disorder [is] selected from the group consisting of osteosarcoma, periodontal disease, fracture healing, osteoarthritis, rheumatoid arthritis, Paget's disease, humoral hypercalcemia, malignancy and osteoporosis comprising administering a compound according to claim 5.~~

10. (Currently Amended) A method according to claim [8] ~~9~~ wherein the bone or mineral disease or disorder is osteoporosis.

11. (Cancelled).

12. (Currently Amended) A method according to claim [7] ~~9~~ wherein the calcilytic compound is co-administered with an anti-resorptive agent.

13. (Original) A method according to claim 12 wherein the anti-resorptive agent is selected from the group consisting of estrogen, 1, 25 (OH)<sub>2</sub> vitamin D3, calcitonin, selective estrogen receptor modulators, vitronectin receptor antagonists, V-H+-ATPase inhibitors, src SH<sub>2</sub> antagonists, bisphosphonates and cathepsin K inhibitors.

14. (Cancelled).